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## **Amendments to the Claims:**

The following Listing of Claims will replace all prior versions, and listings, of the claims in the above-identified application.

## **Listing of Claims**

- 1. (currently amended) An isolated and purified-poly(ADP-ribose) polymerase (PARP) homolog comprising human PARP2 (SEQ ID NO: 2) or a functional equivalent thereof which is at least 95% homologous to or a PARP homolog with at least 85% identity with human PARP2 (SEQ ID NO: 2), which exhibits poly(ADP-ribose)-synthesizing activity, and has an amino acid sequence which
  - a) has a functional NAD<sup>+</sup> binding domain comprising the sequence  $PX_n(S/T)GX_3GKGIYFA \ (SEQ \ ID \ NO:11)$  in which n is an integral value from 1 to 5, and the X radicals are, independently of one another, any amino acid;

and

- b) lacks a zinc finger sequence of the formula  $CX_2CX_mHX_2C \ (SEQ\ ID\ NO:30)$  in which m is an integral value of 28 or 30, and the X radicals are, independently of one another, any amino acid.
- 2. (previously presented) The PARP homolog as claimed in claim 1, wherein the functional NAD<sup>+</sup> binding domain comprises the following sequence:

 $(S/T)XGLR(I/V)XPX_n(S/T)GX_3GKGIYFA (SEQ ID NO:12)$  in which n is an integral value from 1 to 5, and the X radicals are, independently

of one another, any amino acid.

3. (currently amended) The PARP homolog as claimed in claim 1, further comprising the sequence:

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LX<sub>9</sub>NX<sub>2</sub>YX<sub>2</sub>QLLX(D/E)X<sub>10/11</sub>WGRVG (SEQ ID NO: 15)

in which the X radicals are, independently of one another, any amino acid.

- 4-32. (canceled)
- 33. (previously presented) The PARP homolog as claimed in claim 1, wherein the functional NAD<sup>+</sup> binding domain comprises the following sequence:

LLWHG(S/T)X<sub>7</sub>IL(S/T)XGLR(I/V)XPX<sub>n</sub>(S/T)GX<sub>3</sub>GKGIYFAX<sub>3</sub>SKSAXY (SEQ ID NO:13)

in which n is an integral value from 1 to 5, and

the X radicals are, independently of one another, any amino acid.

34. (currently amended) The PARP homolog as claimed in claim 1, further comprising sequence:

AX<sub>3</sub>FXKX<sub>4</sub>KTXNXWX<sub>5</sub>FX<sub>3</sub>PXK (SEQ ID NO:16)

in which the X radicals are, independently of one another, any amino acid.

35. (currently amended) The PARP homolog as claimed in claim 1, further comprising sequence:

XL(I/L)X<sub>2</sub>IX<sub>9</sub>MX<sub>10</sub>PLGKLX<sub>3</sub>QIX<sub>6</sub>L (SEQ ID NO:17)

in which the X radicals are, independently of one another, any amino acid.

36. (currently amended) The PARP homolog as claimed in claim 1, further comprising sequence:

FYTXIPHXFGX<sub>3</sub>PP (SEQ ID NO:18)

in which the X radicals are, independently of one another, any amino acid.

37. (currently amended) The PARP homolog as claimed in claim 1, further comprising sequence:

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## KX<sub>3</sub>LX<sub>2</sub>LXDIEXAX<sub>2</sub>L (SEQ ID NO:19)

in which the X radicals are, independently of one another, any amino acid.

- 38. (currently amended) An isolated poly(ADP-ribose) polymerase (PARP) homolog comprising human PARP2 (SEQ ID NO: 2) or a functional equivalent thereof which is at least 95% homologous to or a PARP homolog having at least 85% identity with human PARP2 (SEQ ID NO: 2), which exhibits poly(ADP-ribose)-synthesizing activity, and has an amino acid sequence which
  - a) has a functional NAD<sup>+</sup> binding domain comprising the sequence  $PX_n(S/T)GX_3GKGIYFA \ (SEQ\ ID\ NO:11)$  in which n is an integral value from 1 to 5, and the X radicals are, independently of one another, any amino acid;

and

b) lacks a zinc finger sequence of the formula

in which m is an integral value of 28 or 30, and the X radicals are, independently of one another, any amino acid

further comprising a leucine zipper-like sequence:

$$(L/V)X_6LX_6LX_6L$$
 (SEQ ID NO: 14)

wherein X radicals are, independently of one another, any amino acid.

39. (currently amended) The PARP homolog as claimed in claim 38, further comprising at least one of the following sequences:

LX<sub>9</sub>NX<sub>2</sub>YX<sub>2</sub>QLLX(D/E)X<sub>10/11</sub>WGRVG (SEQ ID NO: 15), AX<sub>3</sub>FXKX<sub>4</sub>KTXNXWX<sub>5</sub>FX<sub>3</sub>PXK (SEQ ID NO:16),

QXL(I/L)X<sub>2</sub>IX<sub>9</sub>MX<sub>10</sub>PLGKLX<sub>3</sub>QIX<sub>6</sub>L (SEQ ID NO:17),

FYTXIPHXFGX<sub>3</sub>PP (SEQ ID NO:18), and

KX<sub>3</sub>LX<sub>2</sub>LXDIEXAX<sub>2</sub>L (SEQ ID NO:19)

in which the X radicals are, independently of one another, any amino acid.

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40. (currently amended) The PARP homolog as claimed in claim 38, further comprising sequences:

 $LX_9NX_2YX_2QLLX(D/E)X_{10/11}WGRVG$  (SEQ ID NO: 15)

AX<sub>3</sub>FXKX<sub>4</sub>KTXNXWX<sub>5</sub>FX<sub>3</sub>PXK (SEQ ID NO:16),

 $QXL(I/L)X_2IX_9MX_{10}PLGKLX_3QIX_6L$  (SEQ ID NO:17),

FYTXIPHXFGX<sub>3</sub>PP (SEQ ID NO:18), and

KX<sub>3</sub>LX<sub>2</sub>LXDIEXAX<sub>2</sub>L (SEQ ID NO:19)

in which the X radicals are, independently of one another, any amino acid.

41. (currently amended) The PARP homolog as claimed in claim 38, further comprising sequences:

 $LX_9NX_2YX_2QLLX(D/E)X_{10/11}WGRVG$  (SEQ ID NO: 15)

AX<sub>3</sub>FXKX<sub>4</sub>KTXNXWX<sub>5</sub>FX<sub>3</sub>PXK (SEQ ID NO:16),

QXL(I/L)X<sub>2</sub>IX<sub>9</sub>MX<sub>10</sub>PLGKLX<sub>3</sub>QIX<sub>6</sub>L (SEQ ID NO:17),

FYTXIPHXFGX<sub>3</sub>PP (SEQ ID NO:18), and

KX<sub>3</sub>LX<sub>2</sub>LXDIEXAX<sub>2</sub>L (SEQ ID NO:19)

in which the X radicals are, independently of one another, any amino acid, wherein

 $LX_9NX_2YX_2QLLX(D/E)X_{10/11}WGRVG$  (SEQ ID NO:15)

is closest to the N terminus.

42. (currently amended) The PARP homolog as claimed in claim 1, further comprising sequences:

LX<sub>9</sub>NX<sub>2</sub>YX<sub>2</sub>QLLX(D/E)X<sub>10/11</sub>WGRVG (SEQ ID NO: 15)

AX<sub>3</sub>FXKX<sub>4</sub>KTXNXWX<sub>5</sub>FX<sub>3</sub>PXK (SEQ ID NO:16),

 $QXL(I/L)X_2IX_9MX_{10}PLGKLX_3QIX_6L$  (SEQ ID NO:17),

FYTXIPHXFGX<sub>3</sub>PP (SEQ ID NO:18), and

KX<sub>3</sub>LX<sub>2</sub>LXDIEXAX<sub>2</sub>L (SEQ ID NO:19)

in which the X radicals are, independently of one another, any amino acid.

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43. (currently amended) The PARP homolog as claimed in claim 1, further comprising sequences:

LX<sub>9</sub>NX<sub>2</sub>YX<sub>2</sub>QLLX(D/E)X<sub>10/11</sub>WGRVG (SEQ ID NO: 15)

AX<sub>3</sub>FXKX<sub>4</sub>KTXNXWX<sub>5</sub>FX<sub>3</sub>PXK (SEQ ID NO:16),

QXL(I/L)X<sub>2</sub>IX<sub>9</sub>MX<sub>10</sub>PLGKLX<sub>3</sub>QIX<sub>6</sub>L (SEQ ID NO:17),

FYTXIPHXFGX<sub>3</sub>PP (SEQ ID NO:18), and

KX<sub>3</sub>LX<sub>2</sub>LXDIEXAX<sub>2</sub>L (SEQ ID NO:19)

in which the X radicals are, independently of one another, any amino acid, wherein

 $LX_9NX_2YX_2QLLX(D/E)X_{10/11}WGRVG$  (SEQ ID NO:15)

is closest to the N terminus.

44. (currently amended) The PARP homolog as claimed in claim 1, further comprising at least one of the following:

GX<sub>3</sub>LXEVALG (SEQ ID NO: 20),

GX2SX<sub>4</sub>GX<sub>3</sub>PX<sub>a</sub>LXGX<sub>2</sub>V (SEQ ID NO: 21), and

E(Y/F)X<sub>2</sub>YXYX<sub>3</sub>QXYLL (SEQ ID NO: 22)

in which a is 7 to 9 and

X is any amino acid.

45. (currently amended) The PARP homolog as claimed in claim 1, further comprising

GX<sub>3</sub>LXEVALG (SEQ ID NO: 20),

GX<sub>2</sub>SX<sub>4</sub>GX<sub>3</sub>PX<sub>a</sub>LXGX<sub>2</sub>V (SEQ ID NO: 21), and

 $E(Y/F)X_2YX_3QX_4YLL$  (SEQ ID NO: 22)

in which a is 7 to 9 and

X is any amino acid.

46. (currently amended) The PARP homolog as claimed in claim 1, further comprising

GX<sub>3</sub>LXEVALG (SEQ ID NO: 20),

GX<sub>2</sub>SX<sub>4</sub>GX<sub>3</sub>PX<sub>a</sub>LXGX<sub>2</sub>V (SEQ ID NO: 21), and

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 $E(Y/F)X_2YX_3QX_4YLL$  (SEQ ID NO: 22)

in which a is 7 to 9 and

X is any amino acid, wherein

 $E(Y/F)X_2YX_3QX_4YLL$  (SEQ ID NO: 22)

is closest to the C terminus.

- 47. (currently amended) An isolated poly(ADP-ribose) polymerase (PARP) homolog comprising human PARP2 (SEQ ID NO: 2) or a functional equivalent thereof which is at least 95% homologous to or a PARP homologue having at least 85% identity with human PARP2 (SEQ ID NO: 2), which exhibits poly(ADP-ribose)-synthesizing activity, and has an amino acid sequence which
  - a) has a functional NAD<sup>+</sup> binding domain comprising the sequence  $PX_n(S/T)GX_3GKGIYFA \ (SEQ \ ID \ NO:11)$  in which n is an integral value from 1 to 5, and the X radicals are, independently of one another, any amino acid;

and

- b) lacks a zinc finger sequence.
- 48. (previously presented) The PARP homolog as claimed in claim 47, wherein said PARP lacks a zinc finger sequence of the formula

in which m is an integral value of 28 or 30, and

the X radicals are, independently of one another, any amino acid.

49. (previously presented) The PARP homolog as claimed in claim 47, wherein the functional NAD<sup>+</sup> binding domain comprises the following sequence:

 $(S/T)XGLR(I/V)XPX_n(S/T)GX_3GKGIYFA$  (SEQ ID NO:12)

in which n is an integral value from 1 to 5, and

the X radicals are, independently of one another, any amino acid.

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50. (previously presented) The PARP homolog as claimed in claim 47, wherein the functional NAD<sup>+</sup> binding domain comprises the following sequence:

 $LLWHG(S/T)X_7IL(S/T)XGLR(I/V)XPX_n(S/T)GX_3GKGIYFAX_3SKSAXY \ (SEQ\ ID\ NO:13)$ 

in which n is an integral value from 1 to 5, and the X radicals are, independently of one another, any amino acid.

51. (currently amended) The PARP homolog as claimed in claim 47, further comprising a leucine zipper-like sequence:

 $(L/V)X_6LX_6LX_6L$  (SEQ ID NO: 14)

wherein X radicals are, independently of one another, any amino acid.

52. (currently amended) The PARP homolog as claimed in claim 51, further comprising at least one of the following sequences:

LX<sub>9</sub>NX<sub>2</sub>YX<sub>2</sub>QLLX(D/E)X<sub>10/11</sub>WGRVG (SEQ ID NO: 15),

AX<sub>3</sub>FXKX<sub>4</sub>KTXNXWX<sub>5</sub>FX<sub>3</sub>PXK (SEQ ID NO:16),

 $QXL(I/L)X_2IX_9MX_{10}PLGKLX_3QIX_6L$  (SEQ ID NO:17),

FYTXIPHXFGX<sub>3</sub>PP (SEQ ID NO:18), and

KX<sub>3</sub>LX<sub>2</sub>LXDIEXAX<sub>2</sub>L (SEQ ID NO:19)

in which the X radicals are, independently of one another, any amino acid.

53. (currently amended) The PARP homolog as claimed in claim 51, further comprising:

LX<sub>9</sub>NX<sub>2</sub>YX<sub>2</sub>QLLX(D/E)X<sub>10/11</sub>WGRVG (SEQ ID NO: 15),

AX<sub>3</sub>FXKX<sub>4</sub>KTXNXWX<sub>5</sub>FX<sub>3</sub>PXK (SEQ ID NO:16),

QXL(I/L)X<sub>2</sub>IX<sub>9</sub>MX<sub>10</sub>PLGKLX<sub>3</sub>QIX<sub>6</sub>L (SEQ ID NO:17),

FYTXIPHXFGX<sub>3</sub>PP (SEQ ID NO:18), and

KX<sub>3</sub>LX<sub>2</sub>LXDIEXAX<sub>2</sub>L (SEQ ID NO:19)

in which the X radicals are, independently of one another, any amino acid.

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54. (currently amended) The PARP homolog as claimed in claim 51, further comprising:

LX<sub>9</sub>NX<sub>2</sub>YX<sub>2</sub>QLLX(D/E)X<sub>10/11</sub>WGRVG (SEQ ID NO: 15),

AX<sub>3</sub>FXKX<sub>4</sub>KTXNXWX<sub>5</sub>FX<sub>3</sub>PXK (SEQ ID NO:16),

 $QXL(I/L)X_2IX_9MX_{10}PLGKLX_3QIX_6L$  (SEQ ID NO:17),

FYTXIPHXFGX<sub>3</sub>PP (SEQ ID NO:18), and

KX<sub>3</sub>LX<sub>2</sub>LXDIEXAX<sub>2</sub>L (SEQ ID NO:19)

in which the X radicals are, independently of one another, any amino acid, wherein

LX<sub>9</sub>NX<sub>2</sub>YX<sub>2</sub>QLLX(D/E)X<sub>10/11</sub>WGRVG (SEQ ID NO: 15)

is closest to the N terminus.

55. (currently amended) The PARP homolog as claimed in claim 47, further comprising at least one of the following:

GX<sub>3</sub>LXEVALG (SEQ ID NO: 20),

GX<sub>2</sub>SX<sub>4</sub>GX<sub>3</sub>PX<sub>a</sub>LXGX<sub>2</sub>V (SEQ ID NO: 21), and

 $E(Y/F)X_2YX_3QX_4YLL$  (SEQ ID NO: 22)

in which a is 7 to 9 and

X is any amino acid.

56. (currently amended) The PARP homolog as claimed in claim 47, further comprising

GX<sub>3</sub>LXEVALG (SEQ ID NO: 20),

GX<sub>2</sub>SX<sub>4</sub>GX<sub>3</sub>PX<sub>a</sub>LXGX<sub>2</sub>V (SEQ ID NO: 21), and

 $E(Y/F)X_2YX_3QX_4YLL$  (SEQ ID NO: 22)

in which a is 7 to 9 and

X is any amino acid.

57. (currently amended) The PARP homolog as claimed in claim 47, further comprising

GX<sub>3</sub>LXEVALG (SEQ ID NO: 20),

GX<sub>2</sub>SX<sub>4</sub>GX<sub>3</sub>PX<sub>a</sub>LXGX<sub>2</sub>V (SEQ ID NO: 21), and

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 $E(Y/F)X_2YX_3QX_4YLL$  (SEQ ID NO: 22)

in which a is 7 to 9 and

X is any amino acid, wherein

 $E(Y/F)X_2YX_3QX_4YLL$  (SEQ ID NO: 22)

is closest to the C terminus.

58. (currently amended) The PARP homolog as claimed in claim 51, further comprising at least one of the following:

GX<sub>3</sub>LXVALG (SEQ ID NO: 20),

GX2SX<sub>4</sub>GX<sub>3</sub>PX<sub>a</sub>LXGX<sub>2</sub>V (SEQ ID NO: 21), and

 $E(Y/F)X_2YX_3QX_4YLL$  (SEQ ID NO: 22)

in which a is 7 to 9 and

X is any amino acid.

59. (currently amended) The PARP homolog as claimed in claim 51, further comprising

GX<sub>3</sub>LXEVALG (SEQ ID NO: 20),

GX<sub>2</sub>SX<sub>4</sub>GX<sub>3</sub>PX<sub>a</sub>LXGX<sub>2</sub>V (SEQ ID NO: 21), and

 $E(Y/F)X_2YX_3QX_4YLL$  (SEQ ID NO: 22)

in which a is 7 to 9 and

X is any amino acid.

60. (currently amended) The PARP homolog as claimed in claim 51, further comprising

GX<sub>3</sub>LXEVALG (SEQ ID NO: 20),

GX<sub>2</sub>SX<sub>4</sub>GX<sub>3</sub>PX<sub>a</sub>LXGX<sub>2</sub>V (SEQ ID NO: 21), and

 $E(Y/F)X_2YX_3QX_4YLL$  (SEQ ID NO: 22)

in which a is 7 to 9 and

X is any amino acid, wherein

 $E(Y/F)X_2YX_3QX_4YLL$  (SEQ ID NO: 22

is closest to the C terminus.

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61. (new) An isolated and purified-poly(ADP-ribose) polymerase (PARP) homolog comprising human PARP2 (SEQ ID NO: 2).